



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of:
Lan Kluwe

Application No.: 10/692,537

Confirmation No.: 9877

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Art Unit: 1637

For: METHOD FOR THE DETERMINATION OF
DATA FOR THE PREPARATION OF THE
DIAGNOSIS OF PHAKOMATOSIS

Examiner: Kim, Young J.

MS AF
Commissioner for Patents
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Declaration Under 37 C.F.R. §1.132 of Lan Kluwe

Sir:

I, Dr. Lan Kluwe of Hamburg, Germany, hereby declare as follows:

1. I am currently employed by the Laboratory for Tumor Biology and Developmental Disorders, Department of Maxillofacial Surgery, of the University Hospital Eppendorf as Laboratory Head. I have been working and studying in the area of phakomatoses since 1993. I received my doctorate degree in Molecular Biology in 1990. A copy of my resume is attached.
2. I am the inventor on the patent application identified above. I have read an Office Action dated April 17, 2007 relating to this application. I understand that the Examiner is of the view that at the June 27, 2000 filing date of this application, it would have been obvious to determine if an offspring of a patient suffering from a phakomatosis tumor suppressor gene disease is at risk of developing the disease using the method as claimed in my patent application.
3. As of the June 27, 2000 original filing date of this application, even though loss of heterozygosity determinations generally were known, I am not aware that anyone was using this method to determine the risk for offspring. In fact, to my knowledge, at the time my application was filed this testing was done by mutation analysis and occasionally by linkage analysis. Linkage analysis, however, can only be applied to families with at least two affected individuals.
4. I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title XVIII of the United

States Code and that willful false statements may jeopardize the validity of this Application for Patent or any patent issuing thereon.

Lan Kluwe, Ph.D.

Dated: Oct. 15, 2007

Signature: 

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PUBLICATIONS (see detailed list)

First author: 23 (in Mol Hum Genet, Hum Mutat, J Med Genet, FEBS Lett, J Bacteriol, ect.)
Last author: 6
Co-autor: 38

Publikationen

First author

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2. **Lee L**, Imae Y: Role of threonine residue 154 in receptor function of the Tar chemoreceptor in Escherichia coli. J Bacteriol 1990;172:377-382
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11. **Kluwe L**, Friedrich R, Mautner VF: Allelic loss of the NF1 Gene in NF1-associated plexiform neurofibromas. Cytogenet Cancer Genet 1999;110:103-10
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14. **Kluwe L**, Hagel C, Tatagiba M, Thomas S, Stavrou D, Ostertag H, von Deimling A, Mautner VF: Loss of NF1 alleles distinguish sporadic from NF1-associated pilocytic astrocytomas. J Neuropathol Exp Neurol. 2001;60:917-20.
15. **Kluwe L**, Friedrich RE, Tatagiba M, Mautner VF: Presymptomatic diagnosis for children of sporadic neurofibromatosis 2 patients: A method based on tumor analysis. Genet Med. 2002 Jan-Feb;4(1):27-30.
16. **Kluwe L**, Friedrich RE, Korf B, Fahsold R, Mautner VF: NF1 mutations in neurofibromatosis 1 patients with plexiform neurofibromas. Hum Mutat. 2002 Mar;19(3):309.
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- Kluwe L**, Nygren AO, Errami A, Heinrich B, Matthies C, Tatagiba M, Mautner V. 2005. Screening for large mutations of the NF2 gene. Genes Chromosomes Cancer 42(4):384-391.
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Last author

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Co-author

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- 30 Leverkus M, **Kluwe L**, Roll EM, Becker G, Brocker EB, Mautner VF, Hamm H. Multiple unilateral schwannomas: segmental neurofibromatosis type 2 or schwannomatosis? *Br J Dermatol*. 2003 Apr;148(4):804-9.
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